

16805 U.S. PTC

Attorney Docket No. VPI/99-105 DIV

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant(s) : Michael R. Hale et al.

For : INHIBITORS OF ASPARTYL PROTEASE

EXPRESS MAIL CERTIFICATION

"Express Mail" mailing label number: EV 132192199 US Date of Deposit: October 21, 2003

I hereby certify that this transmittal letter and the other papers and fees identified in this transmittal letter as being transmitted herewith are being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 C.F.R. § 1.10 on the date indicated above and are addressed to Mailstop: Patent Application, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

Claire J. Sainti-van Goodman

New York, New York October 21, 2003

Mail Stop: Patent Application Commissioner for Patents P.O. Box 1450

Alexandria, VA 22313-1450

TRANSMITTAL LETTER FOR RULE 53(b) CONTINUING PATENT APPLICATION

Sir:

This is a request for filing a [X] divisional application of pending Application No. 09/591,464, Confirmation No. 2787, filed June 9, 2000.

Transmitted herewith for filing are the

- [X] Specification (367) pages;
- [X] Claims (20 pages);
- [X] Abstract (1 page);
- [X] Preliminary Amendment;
- [X] Information Disclosure Statement and PTO Form-1449
 (w/o references);
- [X] Print EFS Data Sheet;
- [X] 8 sets of copies of Declaration and Power of Attorney with Correction of Inventorship;



- [X] Copy of Consent of Assignee to Correction of Inventorship Under 37 C.F.R. § 1.48;
- [X] Copy of Petition to Correct Inventorship of Non-Provisional Application Under 37 C.F.R. § 1.48(a); and
- [X] 4 sets of copies of the original Declaration and Power of Attorney; for filing in the above-identified patent application.

The enclosed declaration is:

- [] Newly executed (original or copy).
- [X] Copy from a prior application (37 C.F.R. § 1.63(d)).
- [] A signed statement is attached deleting inventors named in the prior application (37 C.F.R. §§ 1.63(d)(2) and 1.33(b)).
- [X] The entire disclosure of the prior application, from which a copy of the declaration is supplied, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.
- [X] The prior application, Application No. 09/591,464, filed June 9, 2000, is assigned to Vertex Pharmaceuticals Incorporated.

The filing fee has been calculated as shown below:

	NUMB:				MBER IRA		RATE	FEE
BASIC FEE							=	\$ 770.00
TOTAL CLAIMS*	31	-	20	=	11	х	\$ 18 =	\$ 198.00
INDEPENDENT CLAIMS**	2	-	3	=	0	х	\$ 86 =	\$ 0.00
FIRST PRESENTATION OF A MULTIPLE DEPENDENT CLAIM + \$290 = \$ 290.00								

^{*} If less than 20, insert 20. TOTAL = \$1,258.00

^{**} If less than 3, insert 3.

- [X] Fee calculation based on claims pending after Preliminary Amendment.
- [X] A check in the amount of \$1,258.00 in payment of the filing fee is transmitted herewith.
- [] Please charge \$____ to Deposit Account No. 06-1075 in payment of the application fee. A duplicate copy of this transmittal letter is transmitted herewith.
- [X] The Director is hereby authorized to charge payment of any additional filing fees required under 37 C.F.R. § 1.16 in connection with the paper(s) transmitted herewith, or credit any overpayment of same, to Deposit Account No. 06-1075. A duplicate copy of this transmittal letter is transmitted herewith.
- [X] The Director is hereby authorized to charge payment of any additional fees that may be due in connection with the paper(s) transmitted herewith, or credit any overpayment of same, to Deposit Account No. 06-1075. A duplicate copy of this transmittal letter is transmitted herewith.

James F. Haley, Jr. (Reg. No. 27,794)

Attorney for Applicants

Min Wang (Reg. No. 51,303)

Kyumin K. Lee (Reg. No. 53,195)

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VPI/99-105

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Examiner

Sonya Wright

Group

1626

Applicant

Michael R. Hale et al.

Application No.

09/591,464

Confirmation No.

2787

Filed

June 9, 2000

Title

INHIBITORS OF ASPARTYL PROTEASE

New York, New York September 23, 2003

Hon. Commissioner for Patents

P.O. Box 1450

Alexandria, Virginia 22313-1450

PETITION TO CORRECT INVENTORSHIP OF NON-PROVISIONAL APPLICATION UNDER 37 C.F.R. § 1.48(a)

Sir:

Pursuant to 37 C.F.R. § 1.48(a), applicants hereby petition to amend the above-identified non-provisional application to correct the inventorship by adding Ronald George Sherrill as an inventor to the application. His address is as follows:

Ronald George Sherrill, a citizen of the United States of America residing at

139 Brannigan Place Cary, NC 27511

The name of Ronald George Sherrill was omitted in this application through error without any deceptive intention on the part of this inventor.

A check in the amount of \$130.00 in payment of the fee set forth in 37 C.F.R. § 1.17(i) is enclosed herewith.

The Director is hereby authorized to charge payment of any additional fee required in connection with this Petition, or to credit any overpayment of same to Deposit Account No. 06-1075. A duplicate copy of this Petition is enclosed.

Respectfully submitted,

James F. Haley, Jr. (Reg. No. 27,794)

Attorney for Applicants

Min Wang (Reg. No. 51,303)

Kyumin K. Lee (Reg. No. 53,195)

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Examiner : Sonya Wright

Group : 1626

Applicants : Michael R. Hale, et al.

Application No. : 09/591,464

Confirmation No. : 2787

Filed : June 9, 2000

FOR : INHIBITORS OF ASPARTYL PROTEASE

New York, New York September 23, 2003

Hon. Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450

CONSENT OF ASSIGNEE TO CORRECTION OF INVENTORSHIP UNDER 37 C.F.R. § 1.48

Sir:

Vertex Pharmaceuticals Incorporated, a corporation organized and existing under the laws of the Commonwealth of Massachusetts and having an office and place of business at 130 Waverly Street, Cambridge Massachusetts 02139-4242, hereby is the assignee of the entire right, title and interest in and to the above-identified patent application by virtue of an assignment from the presently named inventors (a copy of which is enclosed herewith).

As Senior Vice President and General Counsel, I, Kenneth S. Boger, am empowered to act on behalf of Vertex Pharmaceuticals Incorporated, the assignee of the above-identified application.

1

Vertex Pharmaceuticals Incorporated hereby consents to correction of the inventorship of the above-identified patent application by the addition of Ronald George Sherrill as an inventor.

To the best of my knowledge and belief, title is in the name of Vertex Pharmaceuticals Incorporated.

I hereby declare that all statements made herein of my own knowledge are true, and that all statements made on information and belief are believed to be true; and further, that these statements are made with the knowledge that willful false statements, and the like so made, are punishable by fine or imprisonment, or both, under Section 1001, Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

VERTEX PHARMACEUTICALS INCORPORATED

Signed at: COMBRIDGE, MA

Kenneth S. Boger

Šenior Vice President

By:

General Counsel

Date: 9/22/03

&

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Examiner : Not Yet Assigned

Group : Not Yet Assigned

Applicants : Michael R. Hale et al.

Application No.: Not Yet Assigned

Filed : Concurrently Herewith

For : INHIBITORS OF ASPARTYL PROTEASE

New York, New York October 21, 2003

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

STATEMENT UNDER 37 C.F.R. §§ 1.56 AND 1.97(b)

Sir:

Pursuant to 37 C.F.R. §§ 1.56 and 1.97(b), applicants, through their representatives, make of record the following documents:

United States Patents

Inventor	Patent No.	Issue Date
Mohrs et al.	3,743,722	July 3, 1973
Descamps et al.	4,330,542	May 18, 1982
Ryono et al.	4,629,724	December 16, 1986
Martin et al.	5,196,438	March 23, 1993
Kempf et al.	5,354,866	October 11, 1994
Talley et al.	5,622,949	April 22, 1997
Tung et al.	5,723,490	March 3, 1998
Vazquez et al.	5,744,481	April 28, 1998
Amato et al.	5,808,056	September 15, 1998
Vazquez et al.	5,843,946	December 1, 1998

European Patent Applications

0	022	118	published	January 7, 1981
0	181	071	published	March 14, 1986
0	264	795	published	April 27, 1988
0	346	847	published	December 20, 1989
0	364	804	published	April 25, 1990
0	434	365	published	June 26, 1991
0	468	641	published	January 29, 1992
0	486	948	published	May 27, 1992
0	541	168	published	May 12, 1993
0	594	540	published	April 27, 1994

German Patent Application

DE 3542567 published June 5, 1986

Great Britain Patent Application

2,167,759	published	June	4,	1986
2,200,115	published	July	27,	1988

PCT International Patent Applications

WO	90/07329	published	July 12, 1990
WO	91/00725	published	January 24, 1991
WO	91/18866	published	December 12, 1991
WO	92/08688	published	May 29, 1992
WO	92/08698	published	May 29, 1992
WO	92/08699	published	May 29, 1992
WO	92/08700	published	May 29, 1992
WO	92/08701	published	May 29, 1992
WO	92/17176	published	October 15, 1992
WO	93/23368	published	November 25, 1993
WO	93/23388	published	November 25, 1993
WO	93/23379	published	November 25, 1993
WO	94/04491	published	March 3, 1994
WO	94/04492	published	March 3, 1994
WO	94/04493	published	March 3, 1994
WO	94/05639	published	March 17, 1994
WO	94/10134	published	May 11, 1994
WO	94/10136	published	May 11, 1994
WO	94/18192	published	August 18, 1994
WO	94/19322	published	September 1, 1994
WO	95/06030	published	March 2, 1995
WO	95/07269	published	March 16, 1995

WO 95/09843	published April 13, 1995
WO 95/14016	published May 26, 1995
WO 95/32185	published November 30, 1995

Japanese Patent Abstracts

JP	59046252	published	March	15,	1984
JP	59048449	published	March	19,	1984
JP	61071830	published	April	12,	1986

Other Documents

Banker et al., Modern Pharmaceutics, pp. 627-629 (1996).

- R. Bone et al., "X-ray Crystal Structure of the HIV Protease Complex with L-700,417, an Inhibitor with Pseudo C_2 Symmetry", J. Am. Chem. Soc., 113, pp. 9382-84 (1991).
- J.C. Craig et al., "Antiviral Synergy Between Inhibitors of HIV Proteinase and Reverse Transcriptase", Antiviral Chem. and Chemotherapy, 4(3), pp. 161-66 (1990).
- S. Crawford et al., "A Deletion Mutation in the 5' Part of the pol Gene of Moloney Murine Leukemia Virus Blocks Proteolytic Processing of the gag and pol Polyproteins", J. Virol., 53, pp. 899-907 (1985).
- M. Cushman et al., "Delvelopment of Methodology for the Synthesis of Stereochemically Pure Phe ψ [CH₂N]Pro Linkages in HIV Protease Inhibitors", <u>J. Org. Chem.</u>, 56, pp. 4161-67 (1991).
- D.S. Dhanoa et al., "The Synthesis of Potent Macrocyclic Renin Inhibitors", <u>Tetrahedron Lett.</u>, 33, pp. 1725-28 (1992).
- G.B. Dreyer et al., "Hydroxyethylene Isostere Inhibitors of Human Immunodeficiency Virus-1 Protease: Structure-Activity Analysis Using Enzyme Kinetics, X-ray Crystallography, and Infected T-Cell Assays", <u>Biochemistry</u>, 31, pp. 6646-59 (1992).
- G.A. Flynn et al., "An Acyl-Iminium Ion Cyclization Route to a Novel Conformationally Restricted Dipeptide Mimic: Applications to Angiotensin-Converting Enzyme Inhibition", J. Am. Chem. Soc., 109, pp, 7914-15 (1989).

- G. Fontenot et al., "PCR Amplification of HIV-1 Proteinase Sequences Directly from Lab Isolates Allows Determination of Five Conserved Domains", Virology, 190, pp. 1-10 (1992).
- J. Freskos et al., "(Hydroxyethyl)sulfonamide HIV-1 Protease Inhibitors: Identification of the 2-Methylbenzoyl Moiety at P-2", <u>Bio. & Med. Chem. Lett.</u>, 6, pp. 445-450 (1996).
- A. Ghosh et al., "Potent HIV Protease Inhibitors Incorporating High-Affinity P₂-Ligands and (R)- (Hydroxyethylamino) sulfonamide Isostere", <u>Bio. & Med. Chem.</u> Lett., 8, pp. 687-690 (1998).
- E.E. Gilbert, "Recent Developments in Preparative Sulfonation and Sulfation", <u>Synthesis</u>, 1969, pp. 3-10 (1969).
- A. Goldblum, "Modulation of the Affinity of Aspartic Proteases by the Mutated Residues in Active Site Models", FEBS, 261, pp. 241-44 (1990).
- D. Grobelny et al., "Selective Phosphinate Transition-State Analogue Inhibitors of the Protease of Human Immunodeficiency Virus", <u>Biochem. Biophys. Res. Commun.</u>, 169, pp. 1111-16 (1990).
- G.D. Hartman et al., "4-Substituted Thiophene- and Furan-2-sulfonamides as Topical Carbonic Anhydrase Inhibitors", <u>J.</u> Med. Chem., 35, pp. 3822-31 (1992).
- S. J. Hays et al., "Synthesis of cis-4-(Phosphonooxy)-2-piperidinecarboxylic Acid, an N-Methyl-D-aspartate Antagonist", J. Org. Chem., 56, pp. 4984-4086 (1991).
- J.R. Huff, "HIV Protease: A Novel Chemotherapeutic Target for AIDS", <u>Journal of Medicinal Chemistry</u>, 34(8), pp. 2305-14 (1991).
- K.Y. Hui et al., "A Rational Approach in the Search for Potent Inhibitors Against HIV Proteinase", <u>FASEB</u>, 5, pp. 2606-10 (1991).
- Y. Kiso et al., "'O→N Intramolecular Acyl Migration'-type Prodrugs of Tripeptide Inhibitors of HIV Protease", Peptides: Chemistry, Structure and Biology, 61, pp. 157-159 (1996).

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- X. Lin et al., "Enzymic Activities of Two-Chain Pepsinogen, Two-Chain Pepsin, and the Amino-Terminal Lobe of Pepsinogen", J. Biol. Chem., 267(24), pp. 17257-63 (1992).
- K.P. Manfredi et al., "Examination of HIV-1 Protease Secondary Structure Specificity Using Conformationally Constrained Inhibitors", J. Med. Chem., 34, pp. 3395-99 (1991).
- F.R. Marshall, "Computer-Aided Drug Design", <u>Ann. Ref.</u> Pharmacol. Toxicol., 27, pp. 193-213 (1987).
- J.A. Martin, "Recent Advances in the Design of HIV Proteinase Inhibitors", <u>Antiviral Research</u>, 17, pp. 265-78 (1992).
- T.D. Meek et al., "Inhibition of HIV-1 Protease in Infected T-Lymphocytes by Synthetic Peptide Analogues", <u>Nature</u>, 343, pp. 90-92 (1990).
- M. Miller et al., "Structure of Complex of Synthetic HIV-1 Protease with a Substrate-Based Inhibitor at 2.3 Å Resolution", Science, 246, pp. 1149-52 (1989).
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- J.B. Nichols et al., "A Molecular Mechanics Valence Force Field for Sulfonamides Derived by <u>ab</u> <u>initio</u> Methods", <u>J.</u> Phys. Chem., 95, pp. 9803-11 (1991).
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- J.W. Perich et al., "The Synthesis of Multiple O-Phosphoseryl-Containing Peptides via Phenyl Phosphate Protection", J. Org. Chem., 53, pp. 4103-4105 (1988).
- M.S. Plummer et al., "Design of Peptidomimetic Ligands for the pp60^{src} SH2 Domain", <u>Bioorganic & Medicinal Chemistry</u>, 5, pp. 41-47 (1997).
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- M.D. Power et al., "Nucleotide Sequence of SRV-1, a Type D Simian Acquired Immune Deficiency Syndrome Retrovirus" Science, 231, pp. 1567-73 (1986).
- N.A. Roberts, "Rational Design of Peptide-Based HIV Proteinase Inhibitors", Science, 248, pp. 358-61 (1990).
- S. Scharpe et al., "Proteases and Their Inhibitors: Today and Tomorrow", Biochimie, 73, pp. 121-26 (1991).
- S.K. Sharma et al., "Could Angiotensin I Be Produced from a Renin Substrate by the HIV-1 Protease?", Anal. Biochem., 198, pp. 363-67 (1991).
- S. Yamaguchi et al., "Synthesis of HIV Protease Dipeptide Inhibitors and Prodrugs", <u>Peptide Chemistry 1996</u>, pp. 297-300 (1997).

Copies of all the documents listed above were submitted by applicants in the parent U.S. Patent

Application No. 09/591,464; or were cited by the Examiner during prosecution of said parent application. Pursuant to 37 C.F.R. §1.98(d), applicants have not enclosed copies of the listed documents. However, applicants stand ready to provide copies at the Examiner's request.

Applicants respectfully request that the above documents be (1) fully considered by the Examiner during the course of the examination of this application and (2) printed on any patent issuing from this application.

Applicants also request that a copy of the enclosed Form PTO-1449 duly initialed by the Examiner be forwarded to the undersigned with the next communication.

Respectfully submitted,

James F. Haley, Jr. (Reg. No. 27,794)

Attorney for Applicants

Min Wang (Reg. No. 51,303)

Kyumin K. Lee (Reg. 53,195)

Agents for Applicants

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